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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/035,098	12/28/2001	Geoffrey P. Symonds	J&J2084US1	1700

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EXAMINER

SCHNIZER, RICHARD A

ART UNIT	PAPER NUMBER
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1635

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01/23/2008

PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No.	Applicant(s)	
	10/035,098	SYMONDS ET AL.	
	Examiner	Art Unit	
	Richard Schnizer, Ph. D.	1635	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 15 November 2007.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-6,21,22,24-30,32-40,58,60,63-66 and 68-88 is/are pending in the application.
- 4a) Of the above claim(s) 4-6,33-40,58,60,64-66,68 and 79-88 is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-3,21,22,24-30,32,64-66 and 69-78 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☒ The drawing(s) filed on 28 December 2001 is/are: a) ☒ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☒ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☒ All b) ☐ Some * c) ☐ None of:
1. ☒ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date <u>6/22/07;10/12/07;11/20/07</u> . | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

An amendment was received and entered on 11/21/07.

Claims 1-6, 21, 22, 24-30, 32-40, 58, 60, 63-66, and 68-88 remain pending in the application.

Claims 1-3, 21, 22, 24-30, 32, 64-66, and 69-78 read on the elected invention.

Claims 4-6, 33-40, 58, 60, 64-66, 68, and 79-88 stand withdrawn from further consideration pursuant to 37 CFR 1.142(b), as being drawn to a nonelected invention, there being no allowable generic or linking claim. Applicant timely traversed the restriction (election) requirement in the replies filed on 2/14/05, 5/31/06, and 3/5/07.

Priority

Acknowledgment is made of applicant submission of a certified copy of Australian Application No. 3028 as required by 35 U.S.C. 119(b).

Compliance with Sequence Rules

Applicant's amendment placed the application in compliance with the requirements of 37 CFR 1.821 through 1.825.

Claim Rejections - 35 USC § 112

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

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Claims 1-3, 21, 22, 24-30, 32, 64-66, and 69-78 stand rejected under 35 U.S.C. 112, first paragraph, as failing to comply with the enablement requirement. The claim(s) contains subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention.

Claims 1-3, 21, 22, 24-30, 32, 64-66, and 69-78 are drawn to a double stranded RNA complex comprising (a) a first portion which is capable of hybridizing under physiological conditions to at least a portion of an mRNA, (b) a second portion which is capable of hybridizing to the first, wherein said two portions are located on separate molecules, and (c) an additional ribonucleic acid sequence that enhances the ability of dsRNA to alter the expression of the gene encoding the mRNA molecule, wherein the additional ribonucleic acid sequence encodes HIV Tat protein.

The claims do not specify whether the nature of the recited alteration of expression of the recited gene is positive or negative. The specification as a whole and the prior art of record (e.g. Boshier et al (2000), Caplen et al (2000), Fire (1998)) taught the use of double stranded RNA to inhibit gene expression of target genes. There is no evidence or suggestion of record that a dsRNA can be used to augment the expression of any gene to which it has homology, thus one of skill in the art would not believe that the scope of the invention that embraces a positive alteration of gene expression is operable. The specification provides no working example of this embodiment. Accordingly one could not make that embodiment of the invention without undue experimentation.

The art taught that mammalian cells could defend themselves against invasion by dsRNA through several mechanisms including double stranded RNA-activated protein kinase (PKR)-dependent and PKR-independent mechanisms. See Sledz et al (Biochem. Soc. Trans. 32(6): 952-956, 2004, at e.g. Fig. 2 on page 955). For example, when activated by the presence of double stranded RNA, PKR phosphorylates and inactivates elongation initiation factor 2, thereby inhibiting cellular translation. This would be considered to have a global negative effect on gene expression in the affected cell. Other effects include induction of an interferon response subsequent and apoptosis. In other words, dsRNA can inhibit protein expression specifically by an RNAi mechanism, or globally through PKR-dependent and -independent mechanisms. Because Tat is known to inhibit this activity of PKR, it is unclear why one of skill in the art would expect Tat to enhance the ability of a dsRNA to negatively affect expression of a target gene. If the claimed method allowed expression of sufficient Tat to have any effect, one of skill would reasonably expect the effect to be a relief of the translational block. Thus, any target RNAs that had not yet been degraded by the RNAi mechanism would have a greater likelihood of being translated. This in no way correlates with an enhancement of any ability of an dsRNA to alter the expression of a gene encoding a target mRNA. Thus one of skill in the art would not believe that the scope of the invention that embraces a negative alteration of gene expression is operable. The specification provides no working example of this embodiment. Accordingly one could not make that embodiment of the invention without undue experimentation.

In the event that the Examiner has overlooked an enabled embodiment, the following issues remain.

In view of the specification as filed, in order to function as claimed, the Tat protein encoded by the dsRNA must be expressed. However, the claimed dsRNAs do not provide adequate structure to ensure expression of encoded Tat. One of skill in the art appreciates that the RNA encoding Tat would have to be translated to provide Tat protein, and so the RNA encoding Tat would need an operably linked ribosome binding site such as a 5' cap or an internal ribosome entry site, as well as possibly a polyadenylation signal and other translation-/mRNA stability-influencing sequences.

One of skill in the art would not expect a double stranded RNA to be a good substrate for translation because the number of hydrogen bonds formed in the duplex is far larger than the three hydrogen bonds formed between a translation template and a tRNA. Therefore maintenance of dsRNA structure would be thermodynamically more favorable than would be translation of the RNA. Accordingly, one would not expect to achieve efficient expression of Tat, such that it would be unpredictable as to whether or not it could exert its function to the degree required for the invention to function as claimed.

Finally, neither the prior art of record nor the specification provides guidance as to how to simultaneously use a dsRNA for both translation of an encoded sequence, and inhibition of a homologous sequence. Presumably the dsRNA is a substrate for dicer, and/or RISC upon entry into a cell, and these complexes would compete with ribosomes and translation initiation factors for the dsRNA. The specification provides no

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guidance as to how to make the dsRNA such that both translation and dsRNA inhibition can both occur. The instant claims do not limit the strand on which the sequence encoding the Tat protein is located, they set forth no limitation regarding any sequences necessary to ensure translation of the Tat sequence, nor any limitation regarding the relative positions of the Tat sequence and the hybridizing sequences, or the lengths of the strands of the dsRNA. One of skill in the art is left to optimize these variables to obtain simultaneous Tat expression and dsRNA inhibition in the absence of guidance in the prior art or the specification. Accordingly, one of skill would have to perform undue experimentation in order to make the claimed invention.

Response to Arguments

Applicant's arguments filed 11/21/07 have been fully considered but they are not persuasive.

At page 11 of the response Applicant argues that the specification provides details and guidance as to how to structure double stranded RNAs and use them to activate gene expression. Applicant relies for support on the teachings of the specification for dsRNA structure, and on Janowski et al (Nat. Chem. Biol. 2007 Mar:3(3): 166-173) for the use of dsRNAs to activate gene expression. This is unpersuasive. Janowski was published in 2007, well after the filing date of the instant application, and so is not available to Applicant for the purpose of establishing the state of the art at the time of the invention. In any event, Janowski provides substantial evidence supporting the position of the Office. The teachings of Janowski are limited to

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dsRNAs that target promoters. See abstract. The instant claims are not limited to such dsRNAs. In fact, the specification as filed does not even disclose the concept of a dsRNA targeted to a promoter. If this is not the case, then Applicant should point out such support in the specification by page and line number. Janowski also taught that dsRNAs directed against promoters had the effect of silencing transcription from these promoters. See lines 2 and 3 of abstract; and page 166 column 1, second full paragraph, which cites 10 confirmatory references. The instant specification provides no guidance as to how to discriminate between a dsRNA that activates transcription and one that silences transcription. Therefore it is clear that the specification does not provide an enabling disclosure with regard to using dsRNAs targeted to an mRNA to activate the expression of a gene encoding the mRNA.

Also at page 11, Applicant argues that Tat can be used to activate transcription of genes operatively linked to TAR, and that mRNAs comprising both a translatable region and a hairpin siRNA region can be expressed. These arguments are irrelevant because the instant claims require Tat to enhance the ability of recited dsRNA to alter transcription, and as discussed above, the specification does not enable the use of dsRNAs to activate transcription. The specification and prior art teach only how to use dsRNAs, such as those recited in the instant claims, to inhibit gene expression. Because the specification as filed does not teach how to use Tat to enhance the ability of any dsRNA to inhibit gene expression, there is a failure to meet the enablement requirement. For these reasons the rejection is maintained.

Conclusion

No claim is allowed.

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire **THREE MONTHS** from the mailing date of this action. In the event a first reply is filed within **TWO MONTHS** of the mailing date of this final action and the advisory action is not mailed until after the end of the **THREE-MONTH** shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than **SIX MONTHS** from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner(s) should be directed to Richard Schnizer, whose telephone number is 571-272-0762. The examiner can normally be reached Monday through Friday between the hours of 6:00 AM and 3:30 PM. The examiner is off on alternate Fridays, but is sometimes in the office anyway.

If attempts to reach the examiner by telephone are unsuccessful, the Examiner's supervisor, J. Douglas Schultz, can be reached at (571) 272-0763. The official central fax number is 571-273-8300. Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to (571) 272-0547.

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For all other customer support, please call the USPTO Call Center (UCC) at 800-786-9199.

A handwritten signature in black ink, appearing to read 'R. Schnizer', with a stylized, flowing script.

Richard Schnizer, Ph.D.
Primary Examiner
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